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wherein R₁ and R₂ is H, and R₃ is one acyl group containing from 6 to 24 carbon atoms, and where the acyl chains may contain one or more unsaturated bonds, and

b) fatty acids having a formula

where n may be varied between 4 and 22, and where the acyl chain may contain one or more unsaturated bonds.

- 17. The adjuvant composition according to Claim 16, wherein the monoglyceride preparation content is at least 90 %.
- 18. The adjuvant composition according to Claim 16, wherein the monoglyceride preparation content is at least 95%.
- 19. The adjuvant composition according to Claim 16, wherein the acyl chains of the monoglyceride preparations contain 8 to 20 carbon atoms and wherein the acyl chains may contain one or more unsaturated bonds.
- 20. The adjuvant composition according to Claim 16, wherein the acyl chains of the monoglyceride preparations contain 14 to 20 carbon atoms and wherein the acyl chains may contain one or more unsaturated bonds.

- 21. The adjuvant composition according to Claim 16, wherein the acyl chains of the fatty acid contain 8 to 20 carbon atoms, preferably 14 to 20 carbon atoms and wherein the acyl chains may contain one or more unsaturated bonds.
- 22. The adjuvant composition according to Claim 16, wherein the acyl chains of the fatty acid contain 14 to 20 carbon atoms and wherein the acyl chains may contain one or more unsaturated bonds.
- 23. The adjuvant composition according to Claim 16, wherein the antigen comprises an antigen or vaccine that is selected from the group consisting of antigens and vaccines relevant to humans or animals.
- 24. The adjuvant composition according to Claim 23, wherein the animals are marine animals.
- 25. The adjuvant composition according to Claim 16, wherein the composition comprises additional pharmaceutical excipients selected from the group consisting of preservatives and osmotic pressure controlling agents, pH-controlling agents, organic solvents, hydrophobic agents, enzyme inhibitors, water absorbing polymers, surfactants, absorption promoters and anti-oxidative agents.
- 26. The adjuvant composition according to Claim 16, wherein the composition comprises additional adjuvants.
- 27. The adjuvant composition according to Claim 16, wherein the composition is in a form suitable for parenteral or mucosal administration.
- 28. The adjuvant composition according to Claim 27, wherein the composition is in a form suitable for administration to the mucosa of the nose, mouth, vagina, rectum or the intestine.



- 29. The adjuvant composition according to Claim 27, wherein the composition is in a form suitable for administration to the mucosa of the nose.
- 30. A vaccine or antigen composition, containing in 100 g of the final composition:

from 0.01 to 90 g of a antigen or vaccine component from 0.1 to 90 g of a monoglyceride

from 0.1 to 90 g of a fatty acid

from 0.01 to 99 g of water

from 0.01 to 99 g of PBS or salin¢

and optionally one or more additional adjuvarit or excipient.

- 31. A vaccine or antigen composition comprising one or more substances selected from each of the groups
- a) monoglyceride preparations having at least 80 % monoglyceride content and having a formula

wherein R_1 and R_2 is H and R_3 is one acyl group containing from 6 to 24 carbon atoms, and where the acyl chains may contain one or more unsaturated bonds, and

b) fatty acids having a formula



wherein may be varied between 4 and 22, and where the acyl chain may contain one or more unsaturated bonds

in an amount of 0.01 to 15 g/100 ml of the total volume of the composition, and a vaccine or antigen component that is selected from the group consisting of antigens and vaccines relevant to humans or animals, and optionally one or more additional adjuvant or excipient.

- 32. The vaccine or antigen composition according to Claim 31, wherein the monoglyceride preparation content is at least 90%, preferably at least 95%.
- 33. The vaccine or antigen composition according to Claim 32, wherein the monoglyceride preparation content is at least 95%.
- 34. The vaccine or antigen composition according to Claim 31, wherein the acyl chains of the monoglyceride preparations contain 8 to 20 carbon atoms, and wherein the acyl chains may contain one or more unsaturated bonds.
- 35. The vaccine or antigen composition according to Claim 34, wherein the acyl chains of the monoglyceride preparations contain 14 to 20 carbon atoms and where the acyl chains may contain one or more unsaturated bonds.
- 36. The vaccine or antigen composition according to Claim 31, wherein the acyl chains of the fatty acid contain 8 to 20 carbon atoms and where the acyl chains may contain one or more unsaturated bonds.
- 37. The vaccine or antigen composition according to Claim 36, wherein the acyl chains of the fatty acid contain 14 to 20 carbon atoms and where the acyl chains may contain one or more unsaturated bonds.

